

**This is a self-archived version of an original article. This version may differ from the original in pagination and typographic details.**

**Author(s):** van Bijnen, Sam; Muotka, Joonas; Parviainen, Tiina

**Title:** Divergent auditory activation in relation to inhibition task performance in children and adults

**Year:** 2023

**Version:** Published version

**Copyright:** © 2023 The Authors. Human Brain Mapping published by Wiley Periodicals LLC

**Rights:** CC BY 4.0

**Rights url:** <https://creativecommons.org/licenses/by/4.0/>

**Please cite the original version:**

van Bijnen, S., Muotka, J., & Parviainen, T. (2023). Divergent auditory activation in relation to inhibition task performance in children and adults. *Human Brain Mapping*, 44(15), 4972-4985. <https://doi.org/10.1002/hbm.26418>

# Divergent auditory activation in relation to inhibition task performance in children and adults

Sam van Bijnen<sup>1,2</sup>  | Joonas Muotka<sup>1</sup> | Tiina Parviainen<sup>1</sup> 

<sup>1</sup>Centre for Interdisciplinary Brain Research, Department of Psychology, University of Jyväskylä, Jyväskylä, Finland

<sup>2</sup>Faculty of Social and Behavioural Science, Utrecht University, The Netherlands

## Correspondence

Sam van Bijnen, Department of Psychology, University of Jyväskylä, Kärki, Mattilanniemi 6, FI-40014 Jyväskylän yliopisto, Finland.  
Email: [sam.s.vanbijnen@juu.fi](mailto:sam.s.vanbijnen@juu.fi)

## Funding information

Academy of Finland, Grant/Award Number: 311877; H2020 Marie Skłodowska-Curie Actions, Grant/Award Number: 641652

## Abstract

Adults and children show remarkable differences in cortical auditory activation which, in children, have shown relevance for cognitive performance, specifically inhibitory control. However, it has not been tested whether these differences translate to functional differences in response inhibition between adults and children. We recorded auditory responses of adults and school-aged children (6–14 years) using combined magneto- and electroencephalography (M/EEG) during passive listening conditions and an auditory Go/No-go task. The associations between auditory cortical responses and inhibition performance measures diverge between adults and children; while in children the brain–behavior associations are not significant, or stronger responses are beneficial, adults show negative associations between auditory cortical responses and inhibitory performance. Furthermore, we found differences in brain responses between adults and children; the late (~200 ms post stimulation) adult peak activation shifts from auditory to frontomedial areas. In contrast, children show prolonged obligatory responses in the auditory cortex. Together this likely translates to a functional difference between adults and children in the cortical resources for performance consistency in auditory-based cognitive tasks.

## KEYWORDS

auditory processing, cognitive control, development, EEG, MEG, response inhibition

## 1 | INTRODUCTION

Research dedicated to understanding the development of the central auditory system was crucial to reveal the important cortical timings in auditory perception in adults and children. It led to the current understanding that cortical auditory evoked responses change substantially from childhood to adulthood (Paetau et al., 1995; Johnstone et al., 1996; Ponton et al., 2000, 2002; Wunderlich & Cone-Wesson, 2006). These changes coincide with grey and white matter transitions (Moore & Linthicum, 2007) that interact with synaptic signaling to affect the timing and amplitude during development. Indeed, the evoked responses can be interpreted as a measure of cortical network

efficiency, as they rely on trial-by-trial temporal consistency of the cortex to respond to stimuli. Likewise, the timings of the activation pattern are an indicator of (auditory) development (Hämäläinen et al., 2013; Parviainen et al., 2011; van Bijnen et al., 2019). The characteristics of these auditory evoked responses have important links with behavioral skills both in adults and in children (e.g., Johnstone et al., 1996; Näätänen, 1990; Parviainen et al., 2011; van Bijnen et al., 2022), which makes them particularly useful in understanding incremental cognitive competency in human development.

The time-windows of neural activation after auditory stimulation are, however, remarkably different in children and adults. The sequence of these neural responses has originally been characterized

This is an open access article under the terms of the [Creative Commons Attribution](https://creativecommons.org/licenses/by/4.0/) License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

© 2023 The Authors. *Human Brain Mapping* published by Wiley Periodicals LLC.

using electro- and magnetoencephalography recordings at the scalp. The polyphasic adult waveform, as measured by electro- and magnetoencephalography (EEG/MEG), is characterized by intermittent positive deflections at  $\sim 50$  ms (P1) and  $\sim 150$  ms (P2) and a prominent negative deflection at  $\sim 100$  ms (N1). In contrast, children typically show a biphasic response pattern consisting of a positive deflection at  $\sim 100$  ms (P1) and a prolonged negative deflection at  $\sim 250$  ms (N250) (Albrecht et al., 2000; Čeponienė et al., 2002; Orekhova et al., 2013; Picton et al., 1974; Ponton et al., 2000; Ruhnau et al., 2011; Sussman et al., 2008; Takeshita et al., 2002; Wunderlich et al., 2006; Yoshimura et al., 2014). Together, the components reflect neural activity in Heschl's gyri and supratemporal auditory cortices (Shvarts & Mäkelä, 2020). The polarity of the responses depends on recording location (in sensor-level analysis) and anatomy of the neural generators. Changes to the waveform are thought to reflect increased consistency and automation of auditory processing. In the early time-window, the typical adult N1–P2 complex emerges during adolescence (Sussman et al., 2008). However, the activity in the later time-window ( $\sim 250$  ms) attenuates after a certain age. While it is likely a non-linear relationship, it is specific to the child-brain (Parviainen et al., 2011, 2019; Sussman et al., 2008; van Bijnen et al., 2022).

We have previously shown that the later time-window ( $\sim 250$  ms) of auditory activation reflects behaviorally meaningful processes, as it contributes to variance in response inhibition in children even when measured during passive listening conditions (van Bijnen et al., 2022). While there seems to be converging evidence on the behavioral relevance of the later time-window for both auditory-based and more general skills during childhood (Johnstone et al., 1996; van Bijnen et al., 2022), the role of the early time-window responses remains unclear. From one perspective, the early responses ( $< 200$  ms) are exogenous (obligatory) activity and evoked in both adults and children, without any task or attention. However, there are indications that neural processing in this time-window contributes to language and communication ability (Parviainen et al., 2005; Yoshimura et al., 2014) and arousal or attention regulation (Orekhova et al., 2012) in children.

We have suggested that children and adults employ divergent brain mechanisms for a consistent performance in auditory based cognitive tasks. This claim was based on the finding that children relied strongly on the child-specific auditory activation at 250 ms for a consistent performance on (auditory) inhibition tasks (van Bijnen et al., 2022). In contrast, it is well known that adults rely on frontal/medial regions of the cerebral cortex during this time-window for inhibitory and cognitive control processes (Botvinick et al., 2004; Chambers et al., 2009; Falkenstein et al., 1999; Huster et al., 2010; Nieuwenhuis et al., 2003; Smith et al., 2007). Yet, as far as we know, no studies have addressed this discrepancy.

Differences in neural activity pattern complicates comparisons between children and adults in M/EEG studies. Combining M/EEG is favorable because MEG has better signal-to-noise ratio, while EEG provides a better account of deeper and radial sources (Baillet, 2017; Piastra et al., 2021). Therefore, even within adults the different deflections are best picked up by either MEG or EEG depending on

the source locations and orientations (Piastra et al., 2021; Shahin et al., 2007). Indeed, MEG auditory source waveforms vary across individuals depending on the Heschl's gyrus gyrification type (Benner et al., 2017) and the anatomical organization of the auditory cortex (Shaw et al., 2013). In children there is the added difficulty of variability in the developmental stages even within a narrow age-range. Therefore, source locations and orientations might differ between individuals and combined MRI and M/EEG better accounts for these individual differences.

In the present study, we aimed to investigate possible differences in both the spatio-temporal characteristics of activation and the brain-behavior associations between children and adults in the early auditory activation supporting response inhibition. We focused on the early auditory responses that are prevalent in both children and adults and contrasted an auditory Go/No-go task with a passive listening task (Figure 1).

## 2 | MATERIALS AND METHODS

### 2.1 | Participants

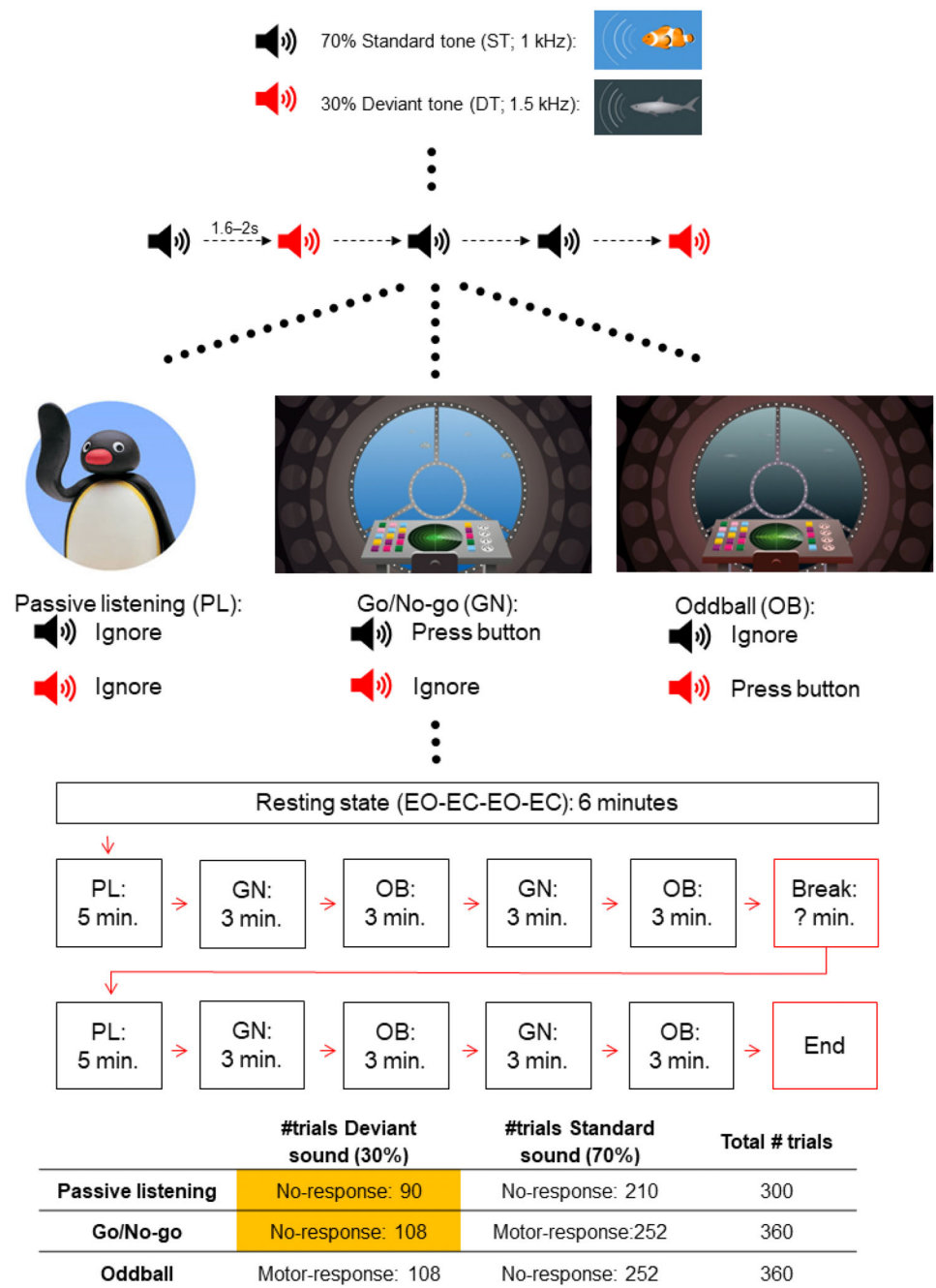
Seventy-eight children that were recruited through schools and the National registry and 16 adults participated in this study. Eleven children and zero adults were excluded due to data quality issues, not finishing the experiment, or structural abnormalities in the MRI. The final dataset consisted of 67 children (36 boys) aged between 6 and 14 years ( $M = 10.2$ ,  $SD = 1.4$ ) and 16 adults (13 women) aged between 20 and 30 years ( $M = 24.8$ ,  $SD = 3.4$ ). All participants had normal hearing as tested with an audiometer. None of the participants had neurological disorders or were on medication affecting the central nervous system. The study was approved by the Ethics Committee of the University of Jyväskylä. An informed consent was obtained from all children and their parents, and the adults in accordance with the Declaration of Helsinki. All participants received compensation for participation (movie ticket or gift card).

The children (but not adults) in this study have previously been described in van Bijnen et al. (2022), which focused on the child-specific N250 activation. Here, we focused on the transient (early) auditory responses that are typically present in both the adults and children.

### 2.2 | Tasks and procedure

The task was presented as a game where the participants were asked to help science by studying and protecting the clownfish population. Studio Dennis Parren ([www.dennisparren.com](http://www.dennisparren.com)) created a visual environment (video) resembling a submarine with a captain giving instructions. All stimuli were controlled by PsychoPy (V3.2) (Peirce et al., 2019) running on a Linux desktop PC. Auditory stimuli were delivered binaurally to the subject through plastic tubes and earpieces using an MEG-compatible hi-fidelity sound system.

**FIGURE 1** Experimental design and procedure. In this study, we specifically focused on the passive listening (PL) Go/No-go (GN) comparison (marked in yellow) to limit the number of statistical tests.



First, we measured resting state activity using two alternating 1.5 min eyes open and closed sessions. Subsequently, the task started with the first passive listening (PL) oddball task. Participants listened to a continuous stream of deviant (DT; 1.5 kHz) and standard (ST; 1 kHz) tones with a 70 ms duration (10 ms rise/fall time) separated with an inter-stimulus interval varying between 1.6 and 2.0 s and were instructed to ignore both tones while they could watch a silent stop-motion video (“Pingu”). After the first PL task participants completed two blocks of alternating Go/No-go (GN) and oddball detection (OB) tasks before the break. Stimuli were identical to the PL task, but participants were asked to either respond to the deviant tone (OB) or the standard tone (GN). After a break, participants completed the

same blocks of the PL, GN and OB tasks as before the break. The complete procedure is shown in Figure 1.

### 2.3 | Behavioral assessment

Cognitive skills were tested on a separate visit. The behavioral tests included subtests of Wechsler Intelligence Scales for Children Third edition (Wechsler, 1999) or Wechsler Adult Intelligence Scale, which was used to evaluate the general level of cognitive skills, and the Stop Signal Task (SST) from the Cambridge Neuropsychological Automated Test Battery (CANTAB).

Of the Wechsler Intelligence scale, the similarities and block design subtests were used to evaluate the general level of verbal and nonverbal reasoning, respectively, and digit span (backward/forward) was used as a measure verbal short-term memory. The coding test and the symbol search task were used to evaluate in general the speed of processing, visuomotor coordination, and attention.

The SST was used to quantify a stop-signal reaction time (SSRT) in adults and children. The SSRT is a behavioral performance measure of response inhibition, or the ability to stop an already initiated motor response which we correlated with the auditory brain activation. In the SST, the participant must respond to an arrow stimulus by selecting one of two options depending on the direction in which the arrow points. The test consists of two parts: in the first part, the participant is first introduced to the test and told to press the left-hand button when they see a left-pointing arrow and the right-hand button when they see a right-pointing arrow. There is one block of 16 trials for the participant to practice this. In the second part, the participant is told to continue pressing the buttons when they see the arrows, but if they hear an auditory signal (a beep), they should withhold their response and not press the button. The task uses a staircase design for the stop signal delay (SSD), allowing the task to adapt to the performance of the participant, narrowing in on the 50% success rate for inhibition. The SST quantifies a stop-signal reaction time (SSRT);

## 2.4 | M/EEG and MRI data analysis

Brain responses were recorded using a 306-channel MEG system and the integrated EEG system (Elekta Neuromag<sup>®</sup> TRIUX™, MEGIN Oy, Helsinki, Finland). M/EEG data were sampled at 1000 Hz and filtered at 0.1–330 Hz. Simultaneous 32-channel EEG and vertical and horizontal electrooculograms (EOG) were recorded with an online reference on the right earlobe. A head position indicator (HPI) continuously monitored the head position in relation to the MEG sensors using five HPI coils, three anatomical landmarks (nasion, left and right preauricular points) and 150 distributed scalp points.

MEG data were first processed and converted to the mean head position with the temporal signal space separation (tSSS) and movement compensation options, implemented in the MaxFilter™ program (version 3.0; MEGIN Oy, Helsinki, Finland). Using MNE-python (Gramfort et al., 2013, 2014) data were low-pass filtered, re-referenced to the average (EEG) and bad channels and data segments were excluded. Epochs were created from –0.2 to 0.8 s relative to stimulus onset and a baseline correction was applied. Epochs from incorrect responses during the task and large MEG signals (>4 pT/cm for gradiometers, >5 pT for magnetometers) were rejected. Independent components representing ocular and/or cardiac artifacts were suppressed with ICA (Hyvärinen & Oja, 2000). Epochs were checked manually and with *autoreject* (EEG) (Jas et al., 2017) to repair or exclude bad epochs.

We used cortically-constrained, depth-weighted ( $p = .8$ ) L2 minimum norm estimate (Hämäläinen & Ilmoniemi, 1994) with a loose orientation constraint (0.2) to characterize the source currents.

T1- and T2-weighted 3D spin-echo MRI images were collected with a 1.5 T scanner (GoldSeal Signa HDxt, General Electric, Milwaukee, WI, USA) using a standard head coil and with the following parameters: TR/TE = 540/10 ms, flip angle = 90°, matrix size = 256 × 256, slice thickness = 1.2 mm, sagittal orientation. The cortical surface was constructed from the individual MRIs with the Freesurfer software (RRID: SCR\_001847, Martinos Center for Biomedical Imaging, <http://freesurfer.net>; Dale et al., 1999; Fischl, Sereno, & Dale, 1999; Fischl, Sereno, Tootell, et al., 1999). Next, the M/EEG data were registered to the individual structural data with MNE coregistration using the anatomical landmarks, digitized EEG electrodes and additional scalp points. The forward solution for the source space was constructed using a three-layer BEM with the following conductivity values for brain/CSF, skull, and scalp: 0.3, 0.006 and 0.3 for adults and 0.33, 0.0132 and 0.33 for children. The noise covariance matrix was calculated from the individual epochs 200-ms pre-stimulus baseline, using a cross validation method implemented in MNE. The MEG and EEG data were combined into a single inverse solution by a whitening transformation using the covariance matrix (Engemann & Gramfort, 2015). Therefore, the inverse solvers estimating the source time courses are informed by both the MEG and the EEG signals.

The final source waveforms were computed as the mean value within the transverse temporal gyrus (label 30 in the Desikan-Killiany Atlas; Desikan et al., 2006). The time-window for the extraction of amplitude values were based on the grand averages (Figure 3) where the time-windows for the main peaks corresponded to those reported in earlier literature in children and adults (Albrecht et al., 2000; Orekhova et al., 2013; Picton et al., 1974; Ponton et al., 2000; Yoshimura et al., 2014). In children, we extracted the maximum value between 76 and 104 ms (P1), minimum value between 108 and 140 ms (N1) and the difference between N1 and max value between 148 and 200 ms (P2). In adults we extracted the same P1–N1–P2 responses in different time-windows; 40–76 ms, 92–120 ms and 140–200 ms respectively.

## 2.5 | Statistical analysis

We focused on the adult and child differences in the Passive Listening (PL) versus Go/No-Go task (Figure 1). Task requirements in the Go/No-Go task specifically engaged inhibition. Moreover, our earlier (van Bijnen et al., 2022) study indicated that although both oddball and go/no-go comparison both yielded behaviorally significant activation, the Go/No-go showed stronger correlations. To limit unnecessary statistical testing, we therefore analyzed the deviant tones of these tasks. Importantly, between the tasks, the stimuli (DT), probability (30%) and motor response (none) were identical. Auditory responses (P1–N1–P2) were analyzed separately; models contained one of the brain responses (P1, N1 or P2) as dependent variable at a time. Two within-subject independent (hemisphere (left, right) and task (passive, no-go)) and one between-subject variable (group; children vs. adults) were included in the model. Models were estimated by using Multigroup analysis with Mplus statistical package (Version

8.4) and using a full information maximum likelihood (FIML) estimation method with robust standard errors (MLR). Multigroup models test models in two or more discrete groups. Equality constraints across groups are used to conduct nested tests using likelihood ratio comparisons between a model with certain parameters constrained to be equal and a model with those same parameters freely estimated (allowed to differ) across the groups. Multigroup FIML estimation methods do not have the same equality of (co)variances assumptions and are therefore not hampered by unequal sample sizes. This, however, does not negate the loss of statistical power with smaller sample sizes. All available data were used in the analyses and missing data were assumed to be Missing at Random (MAR) (Muthén & Muthén, 2012). Interactions and main effects were estimated by using additional parameters of model.

Subsequently, we used (partial) correlations (corrected for age in children) to test for relevant brain-behavior associations of the auditory responses. We included the following behavioral performance measures: intra-individual coefficient of variation (ICV; calculated as SDRT/mean RT) (Stuss et al., 2003), response accuracy (RA; calculated as the square root of the error%) and the stop-signal reaction time (SSRT). Correlational analysis was performed with SPSS statistics 25.

### 3 | RESULTS

#### 3.1 | Descriptive statistics of cognitive skills and behavioral performance

Descriptive statistics of the children's performance during the M/EEG experiment and their cognitive skills as per the behavioral assessment session are presented in Table 1. The subtests of the Wechsler Intelligence Scale were included to examine whether participants are within

typical range of cognitive performance and were not associated with the brain responses of children and adults ( $p > .05$ ). Figure 2 shows the relevant age-behavior relationships.

#### 3.2 | Age related differences in the auditory evoked responses

Figure 3 shows the measured neuromagnetic responses to the tones in the passive listening and Go/No-go task at MEG sensor level (gradiometers). For visualization purposes, groups were separated by age (<10 years old, >10 years old and adults). The activation pattern in younger children indicate three separate peaks at ~85ms, at ~120ms and at ~250 ms. The waveforms get progressively more dissociated with age (Figure 3), possibly signifying temporal overlap between components in sensor-level data, especially in the youngest group. The analogous source-level waveforms of the different age groups that depict the possible developmental trajectory of the responses in the auditory cortex are included in Figure S1. Furthermore, responses appear different in the two hemispheres in children: with the P1m predominantly showing in the left hemisphere, the N1m predominantly showing in the right hemisphere and the late N250m showing bilaterally. In contrast, the activation peaks in adults occur somewhat earlier, at ~60 ms and at ~110 ms, and the peak at ~250 ms is clearly diminished. Adults also show less hemispheric differences than children.

#### 3.3 | Differences in child and adult auditory responses

The localization of auditory activation in children indicates that the peaks across the entire timeline of activation all reflect cortical

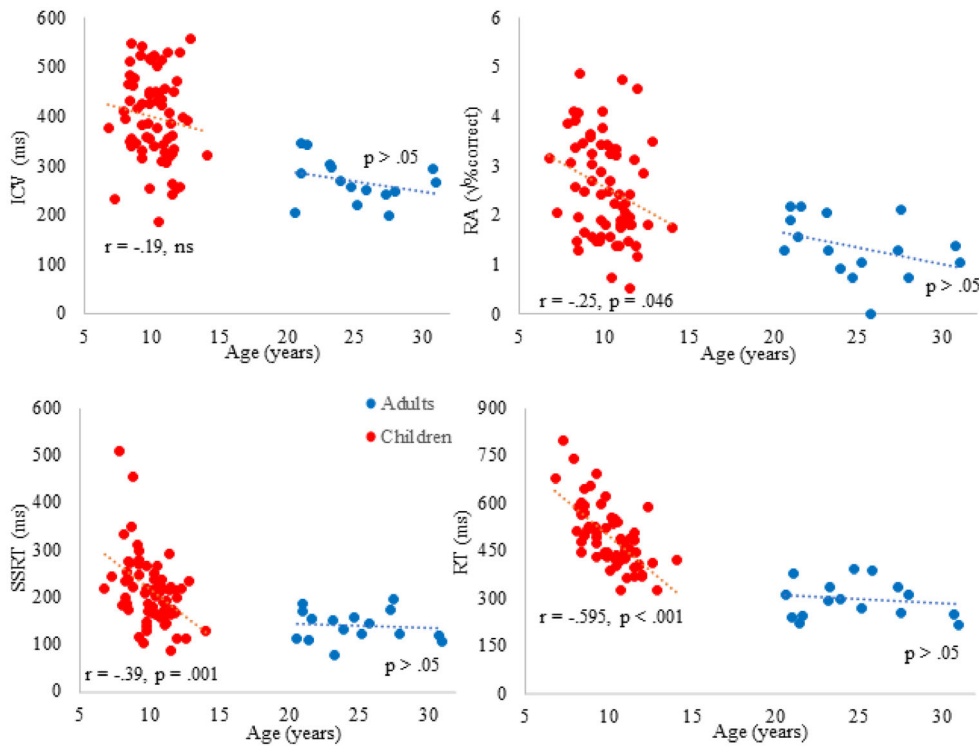
**TABLE 1** Mean, standard deviation (SD) and range of behavioral performance measures.

	Children			Adults		
	Mean	SD	Range	Mean	SD	Range
Age (years)	10.17	1.44	6–14	24.78	3.38	20–30
M/EEG experiment						
GN RT (ms)	484.20	82.74	328–693	298.50	57.5	221–395
GN ICV	0.4	0.09	0.19–0.56	0.27	0.05	0.2–0.35
GN RA (% correct)	2.54	1	0.53–4.87	1.36	0.62	0–2.17
Behavioral assessment						
SSRT (ms)	205.94	56.20	87–351	140.81	32.62	80–198
Digit span <sup>a,b</sup>	10.55	2.65	5–17	10.38	2.36	8–17
Symbol search <sup>a</sup>	12	2.58	5–18	11.06	3.25	4–16
Coding <sup>a</sup>	10.88	2.98	4–19	12.25	1.71	9–16
Block design <sup>a</sup>	11.61	2.97	4–17	12.94	4.46	6–19
Similarities <sup>a</sup>	10.39	2.63	2–16	10.31	2.71	7–16

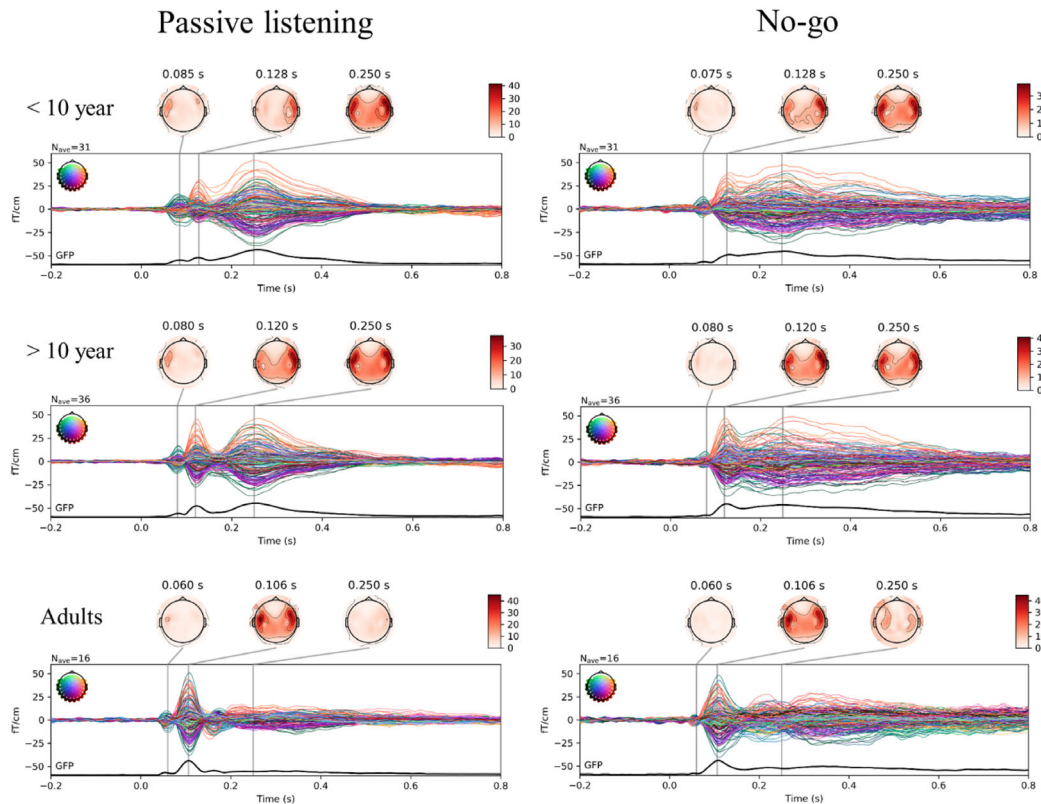
Note: Reaction times (RT), intra-individual coefficient of variation (ICV) and response accuracy (RA) gathered from the Go/No-go task (GN) and the Oddball task (OB). Stop-signal reaction time (SSRT) was gathered from the stop-signal task during the behavioral assessment.

<sup>a</sup>Standardized scores.

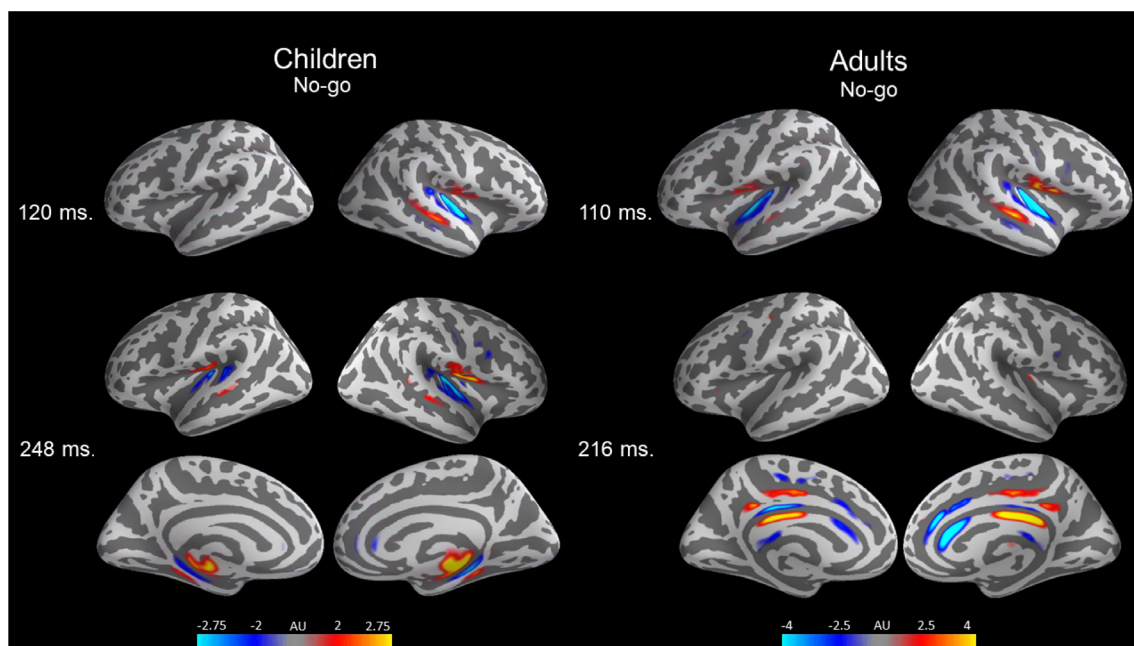
<sup>b</sup>Digit span forward and backward.



**FIGURE 2** Age effects in children (red) and adults (blue) on behavioral data: intra-individual coefficient of variation (ICV), response accuracy (RA), stop-signal reaction time (SSRT) and mean reaction time (RT).



**FIGURE 3** Sensor-level developmental (age) differences in auditory brain responses in the passive listening (left) and Go/No-go (right) task as measured by the MEG gradiometers. Groups divided for illustration purposes between children younger than 10 (top), older than 10 (middle) and adults (bottom).



**FIGURE 4** Grand average 3D visualization of the No-go (deviant tone) M/EEG combined source estimates for all children (right) and adults (left). 3D-plots are presented for the two most prominent time-windows of activation in children (120 and 248 ms) and adults (110 and 216 ms).

currents in the temporal regions irrespective of task and time-window (Figures 4 and 5). In contrast, in adults the early peak at 100 ms reflects activation in the temporal regions and the later activation at ~200–300 ms reflects activation in the medial regions of the cerebral cortex (e.g., cingulate cortex; Figure 4).

The activation pattern at ~250 ms in the auditory cortex is unique to the child brain (Figures 4 and 5) in our data. Consequently, the activation pattern in children versus adults in the 250-ms time-window reflects different brain regions, their strength is not directly comparable. Therefore, we did not directly contrast adults and children for this activation pattern. The behavioral relevance of this activation pattern in children is discussed elsewhere (van Bijnen et al., 2022). Due to the spatiotemporal different nature of the activation at 250 ms, the statistical analysis was limited to the earlier (50–200 ms) auditory cortex responses (P1–N1–P2).

### 3.4 | Passive versus Go/No-go

#### 3.4.1 | P1

In children both task ( $p < .001$ ) and hemisphere ( $p = .001$ ) significantly affected the amplitude of the P1 (Figure 6). The task effect was similar in the two hemispheres as the task  $\times$  hemisphere interaction was not significant ( $p = .301$ ). More specifically, the P1 in the PL task was stronger compared to the P1 in GN task in the left ( $M = 7.69$ ,  $SD = 8.13$  vs.  $M = 4.38$ ,  $SD = 7.86$ ) and right hemisphere ( $M = 3.08$ ,  $SD = 5.84$  vs.  $M = 0.01$ ,  $SD = 7.61$ ). The P1 in children was significantly stronger in the left, compared to the right hemisphere in both

the PL and GN task ( $p < .001$ ) (Figure 6). Finally, there was no main effect of group on the P1 ( $p = .296$ ).

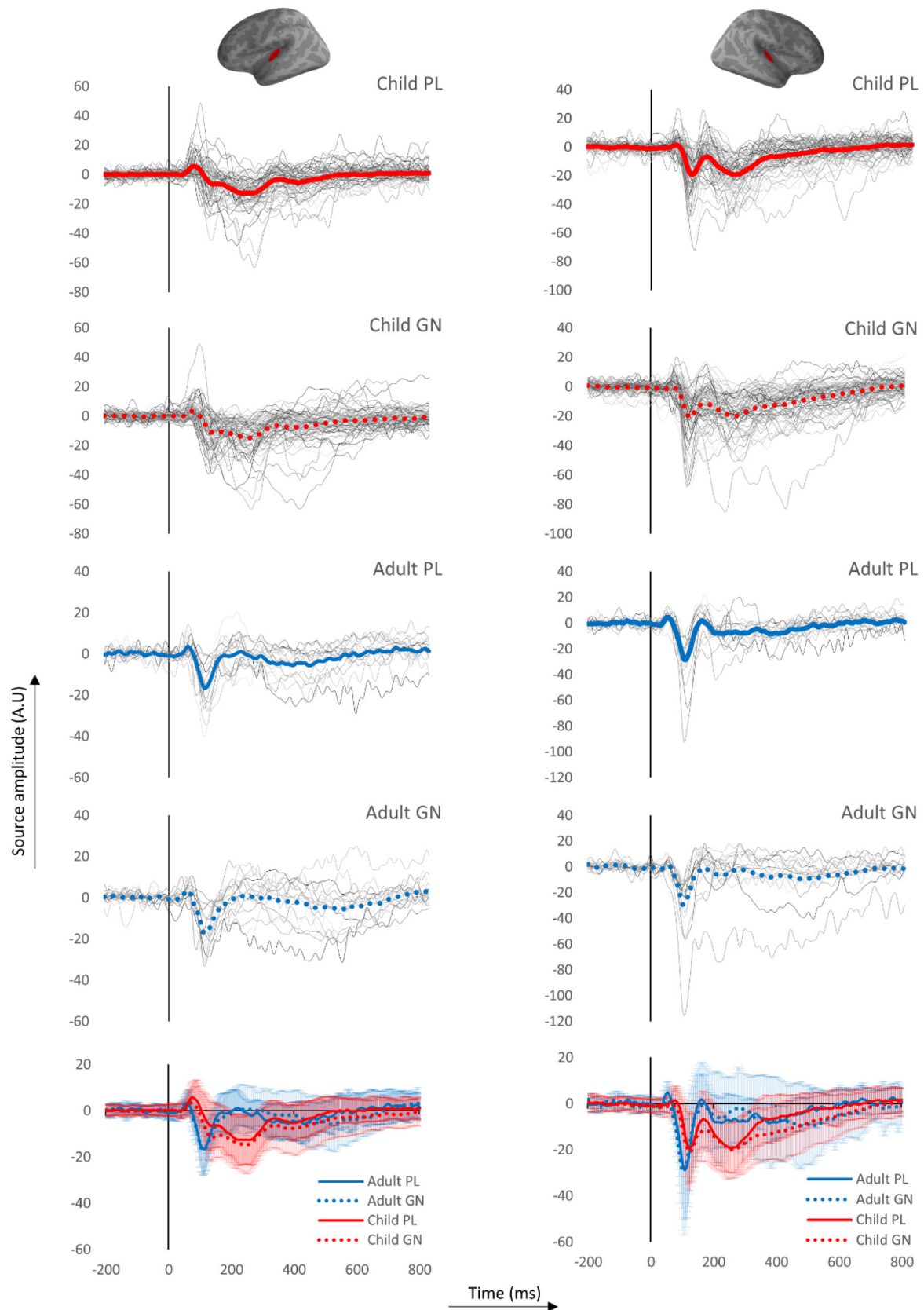
Adults showed the same task effect as children, but no hemisphere effect since the group  $\times$  task interaction was not significant ( $p = .163$ ) but the group  $\times$  hemisphere interaction was significant ( $p = .012$ ). The task effect was similar in the two hemispheres as the task  $\times$  hemisphere interaction was not significant ( $p = .301$ ) and the group  $\times$  task  $\times$  hemisphere interactions ( $p = .204$ ) were not significant. Like the child group, the P1 in the PL task was stronger compared to the P1 in GN task in the left ( $M = 5.59$ ,  $SD = 4.44$  vs.  $M = 4.7$ ,  $SD = 3.97$ ) and right hemisphere ( $M = 6.13$ ,  $SD = 4.87$  vs.  $M = 2.91$ ,  $SD = 4.02$ ) (Figure 6).

The P1 amplitude did not consistently correlate with age in children, with only the No-go P1 in the right hemisphere reaching significance ( $r = -.28$ ,  $p = .02$ ). No significant correlations were found between the P1 in children and their behavioral performance. In adults however, stronger No-go P1 in the left hemisphere was related to a higher ICV (more variability) ( $r = .69$ ,  $p = .003$ ). Table 2 shows the correlation matrix and Figure 7 the associated scatterplot. Figure S2 depicts the scatterplot with separated age groups in children (<10-year-old and >10-year-old).

#### 3.4.2 | N1

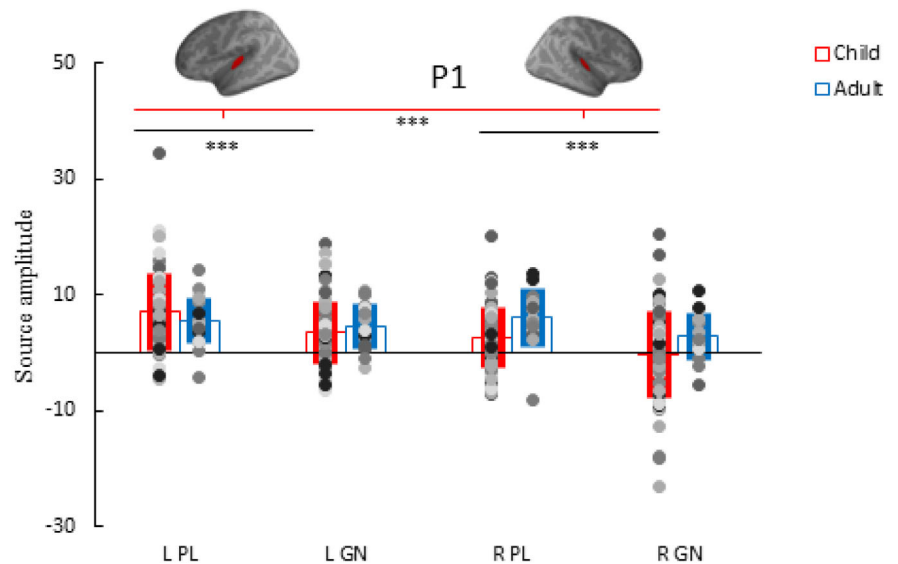
Children and adults showed similar N1 main effects as the N1 showed no significant interactions between group, task, and hemisphere ( $p > .05$ ). There were main effects of group and hemisphere, but the task did not influence N1 strength ( $p > .05$ ). Adults showed stronger N1 responses compared to children ( $p = .044$ ) and the right





**FIGURE 5** M/EEG combined Source estimates in the left and right transverse temporal gyrus (red area) of adults (blue lines) and children (red lines). Waveforms are an average of the entire area. Figures depict the passive (solid lines) and inhibition (No-go) (dotted lines) waveforms in the left (top) and right (bottom) hemisphere. Shaded areas around the waveform represent the standard deviation (SD).

**FIGURE 6** P1 amplitude individual data points, group average and standard deviation for the conditions: passive listening (PL) deviant tone and Go/No-go (GN) deviant tone in the left (L) and right (R) hemisphere of adults (blue) and children (red). \*\*\* $p < .001$ .



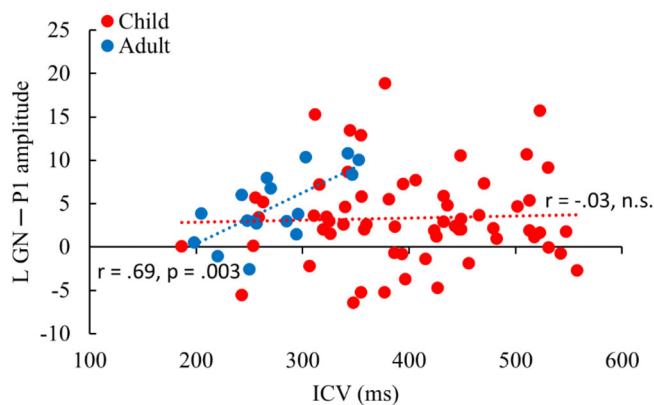
**TABLE 2** Partial (correcting for age in children only) correlation between the P1 amplitude and behavioral measures.

	Child P1			Adult P1		
	ICV	RA	SSRT	ICV	RA	SSRT
L PL	-.052	.019	-.010	.117	-.024	-.116
R PL	.146	.180	-.072	.308	.164	-.047
L GN	-.034	-.005	-.105	<b>.689**</b>	.284	.270
R GN	.206	.189	-.020	-.022	-.162	-.191

Note: Significant correlations marked in bold.

Abbreviations: ICV, intra-individual coefficient of variability; RA, response accuracy; SSRT, stop signal reaction time.

\*\* $p < .01$ .



**FIGURE 7** Scatterplot of the left hemisphere P1 response to the No-go tone and the intraindividual coefficient of variability (ICV) of the Go/No-go (GN) task of children (red) and adults (blue).

hemisphere N1 responses were stronger compared to left hemisphere N1 responses in both tasks and groups ( $p < .001$ ) (Figure 8).

The N1 did not consistently correlate with age in children, with only the No-go N1 in the right hemisphere reaching significance

( $r = -.26$ ,  $p = .035$ ). In children, a stronger left hemisphere N1 response in the PL task is associated with lower ICV ( $r = .302$ ,  $p = .03$ ). In contrast, adults show opposite associations with a stronger left hemisphere N1 response associated with higher ICV (Table 3) and poorer performance in both the PL ( $r = -.701$ ,  $p = .003$ ;  $r = -.67$ ,  $p = .004$ , respectively) and GN ( $r = -.612$ ,  $p = .012$ ;  $r = -.793$ ,  $p < .001$ , respectively) task. Table 3 shows the correlation matrix and Figure 9 the associated scatterplots. Figure S3 depicts the scatterplots with separated age groups in children (<10-year-old and >10-year-old).

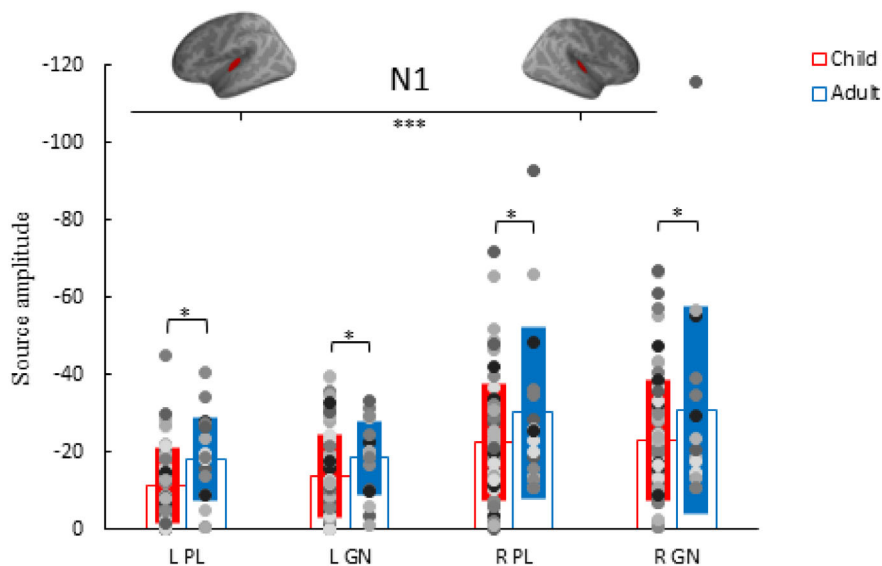
### 3.4.3 | P2

Like the N1, the P2 showed no significant interaction between group, task and hemisphere ( $p > .05$ ). There were main effects of group and hemisphere, but task did not significantly affect P2 amplitudes ( $p > .05$ ). Adults showed stronger P2 responses compared to children ( $p < .001$ ) and the right hemisphere P2 responses were stronger compared to left hemisphere N1m responses in both tasks and groups ( $p < .001$ ) (Figure 10). The P2 showed consistent age effects with the correlation coefficient ranging between .3 and .351 ( $p \leq .014$ ).

The (partial) correlation revealed that, in children, a stronger No-go P2 in the right hemisphere is associated with lower SSRTs ( $r = -.331$ ,  $p = .008$ ). In adults, left hemisphere No-go P2 responses were associated with higher SSRT ( $r = .55$ ,  $p = .027$ ), higher ICV ( $r = .538$ ,  $p = .031$ ) and worse response accuracy ( $r = .535$ ,  $p = .033$ ). Table 4 shows the correlation matrix and Figure 11 the associated scatterplots. Figure S4 depicts the scatterplots with separated age groups in children (<10-year-old and >10-year-old).

## 4 | DISCUSSION

In this study we found key differences in the auditory processing of adults and children. First, children rely on auditory areas for an



**FIGURE 8** N1 amplitude individual data points (dots), group average (empty bar) and standard deviation (solid bar) for the conditions: passive listening (PL) deviant tone and Go/No-go (GN) deviant tone in the left (L) and right (R) hemisphere of adults (blue) and children (red). \* $p < .05$ , \*\*\* $p < .001$ .

**TABLE 3** Partial (correcting for age in children only) correlation between the N1 amplitudes and behavioral measures.

	Child N1			Adult N1		
	ICV	RA	SSRT	ICV	RA	SSRT
L PL	<b>.302*</b>	.249	.145	<b>-.701**</b>	<b>-.670**</b>	-.394
R PL	.082	.101	.092	.067	-.148	.062
L GN	.222	.231	.087	<b>-.612*</b>	<b>-.793***</b>	-.448
R GN	.102	.107	.108	-.062	-.290	.017

Note: Significant correlations marked in bold.

Abbreviations: ICV, intra-individual coefficient of variability; RA, response accuracy; SSRT, stop signal reaction time.

\* $p < .05$ . \*\* $p < .01$ . \*\*\* $p < .001$ .

**TABLE 4** Partial (correcting for age in children only) correlation between the P2 amplitudes and behavioral measures.

	Child P2			Adult P2		
	ICV	RA	SSRT	ICV	RA	SSRT
L PL	-.122	-.189	-.051	<b>.647**</b>	.434	.315
R PL	-.084	-.002	<b>-.278*</b>	.053	.197	.157
L GN	-.045	-.131	-.107	<b>.538*</b>	<b>.535*</b>	<b>.550*</b>
R GN	-.128	-.048	<b>-.331**</b>	.000	.162	-.090

Note: Significant correlations marked in bold.

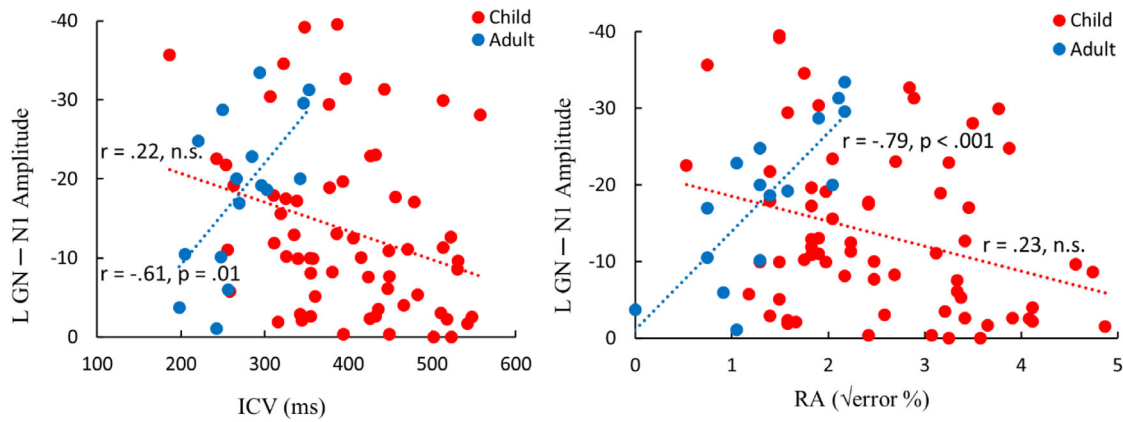
Abbreviations: ICV, intra-individual coefficient of variability; RA, response accuracy; SSRT, stop signal reaction time.

\* $p < .05$ . \*\* $p < .01$ .

extended period to process auditory information during a Go/No-go task compared to adults; the adult activation pattern shifts at ~200 ms from auditory to medial regions of the cerebral cortex that are implicated in cognitive control processing, whereas children show prolonged obligatory responses in the auditory cortex. This indicates a difference in cognitive control processing between adults and children. Second, the associations between the early auditory cortical

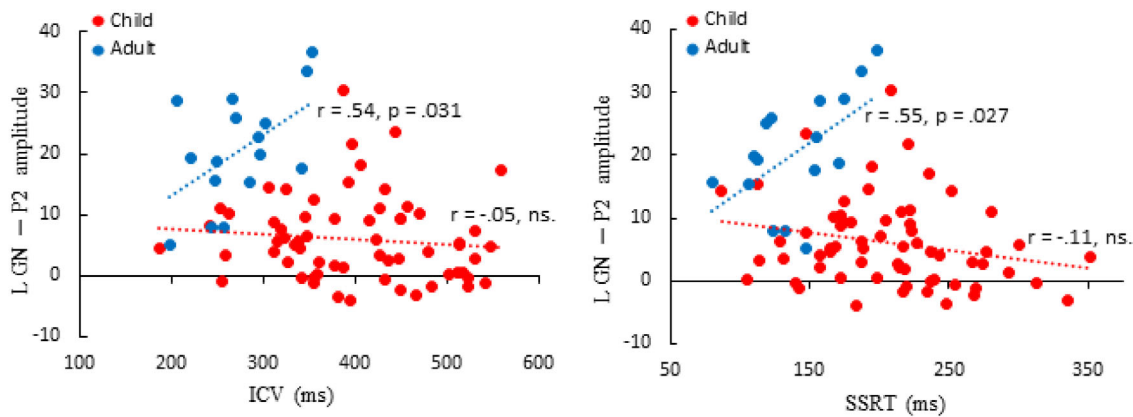
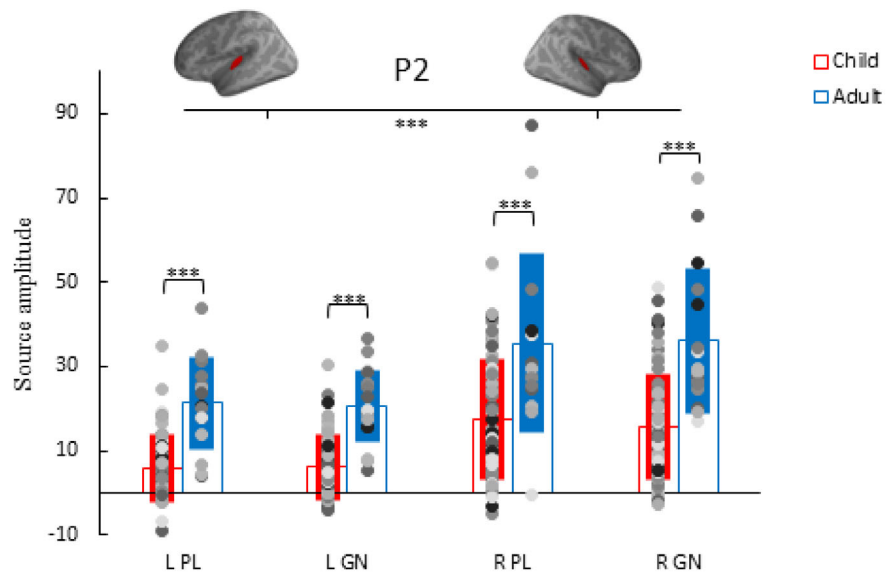
responses and inhibition performance measures diverge between adults and children; while in children the brain-behavior associations are not significant, or stronger responses are beneficial, adults show negative associations between auditory cortical responses and inhibitory performance. Together this likely translates to a functional difference between adults and children in the cortical resources for performance consistency in auditory-based cognitive control tasks.

The developmental changes and age differences (Figure 3; Figure S1) that have been indicated to reflect more efficient (or automated) auditory processing are a more pronounced N1 and in general a gradual temporal dissociation, resulting in clearer separation, of the earlier responses (P1-N1-P2) and an attenuation of the later N2 response (Albrecht et al., 2000; Čeponienė et al., 2002; Ponton et al., 2000; Sussman et al., 2008; Takeshita et al., 2002; Wunderlich & Cone-Wesson, 2006). The age-related dynamics, specifically the decrease in P1 at the early stages of development and a subsequent increase in N1, are in line with the present findings and are together likely to reflect maturational differences in the cortical circuitry (Orekhova et al., 2012). Notably, deep layers (lower III to IV) in the auditory cortex mature earlier, between 6 months and 5 years of age, compared to the superficial layers (upper III and II) that mature somewhere between 6 and 12 years (Eggermont & Ponton, 2003; Moore & Guan, 2001; Moore & Linthicum Jr, 2007; Ponton et al., 1999). As we studied older children, their cortical P1 generators arguably closely reflect that of adults, unlike the neural generators of the N1 that are still developing in our sample of children. Accordingly, we found no age or group effects of the P1 but did for the N1 and P2. A similar explanation could account for our task effect in the P1 that showed smaller amplitudes to the No-go compared to the passive tones in both adults and children, but not in the N1 and P2. Indeed, highlighting the different neural generators of the P1 and N1/P2, a study modelling the adult auditory responses reported that an initial excitatory thalamocortical feedforward drive to layer II/III and V, via layer IV induced the P1. In contrast, a cortico-cortical feedback input to supragranular layers with a subsequent second feedforward input



**FIGURE 9** Scatterplot of the left hemisphere N1 responses to the No-go tone and the behavioral performance measures of the Go/No-go (GN) task: intraindividual coefficient of variability (ICV; left) and response accuracy (RA; right) of children (red) and adults (blue).

**FIGURE 10** P2 amplitude individual data points (dots), group average (empty bar) and standard deviation (solid bar) for the conditions: passive listening (PL) deviant tone and Go/No-go (GN) deviant tone in the left (L) and right (R) hemisphere of adults (blue) and children (red). \*\*\* $p < .001$ .



**FIGURE 11** Scatterplot of the left hemisphere P2 responses to the No-go tone and the behavioral performance measures of the Go/No-go (GN) task: intraindividual coefficient of variability (ICV; left) and stop-signal reaction times (SSRT; right) of children (red) and adults (blue).

induced the N1 and P2 respectively (Kohl et al., 2022). Apparently, the necessity to inhibit a response lowers the thalamocortical input that is associated with the P1.

In addition to the amplitude and latency changes to the auditory responses, their function also changes; children seem to rely more strongly on sensory activation in active (inhibition) tasks. We show

that, while the adult's early auditory components (i.e., P1–N1–P2) are all negatively related to inhibitory performance, the early responses in children are not, or are (weakly) positively related. Instead, the child-specific auditory activation pattern at ~250 ms post-stimulation (N2) in the left hemisphere is reported to be strongly (positively) related to inhibitory performance (Johnstone et al., 1996; van Bijnen et al., 2022).

Earlier studies have highlighted the discrepancies between the early (P1–N1–P2) and the later, obligatory, (N2) response pattern in children, calling it “the additional process” (Johnstone et al., 1996) and suggesting it reflects the child's wider range of attentional focus, resulting in similar neural response patterns in distinct situations (e.g., active vs. passive) while still behaviorally relevant (Johnstone et al., 1996; van Bijnen et al., 2022). Our data supports this claim as we were unable to falsify that the early response amplitudes are not behaviorally relevant for inhibitory performance in children. Thus, the child-specific, obligatory, N2 response seems to be unique in its behavioral relevance compared to the earlier obligatory auditory responses in children. Another possibility is that the early auditory child responses are also beneficial for task performance, but perhaps the child and adult brain–behavior associations are diametrically opposed. This would complicate matters as the brain–behavior associations possibly “flip” during development and this could limit the ability to detect brain–behavior associations in the P1–N1–P2 complex in children.

Our data are in line with previous studies on the adult neurobiological mechanisms of response inhibition and shows it is remarkably different in children in auditory based tasks. In adults, response inhibition is supported by a broad frontoparietal network (Puiu et al., 2020; Weiss & Luciana, 2022) and similarly the midcingulate cortex is reported to be a major neural generator of the adult N2 in active tasks (Huster et al., 2010). However, we show that, in children, the major source of activation in this time-window during No-go trials is in the auditory cortex. It suggests that this auditory activation pattern in children becomes obsolete as the brain becomes more efficient in discriminating auditory stimuli and determining their behavioral relevance. The child brain is not merely an “miniature” adult brain but the mechanisms that govern inhibitory performance in children are functionally distinct from adults. The maturational changes in the auditory response coincide with improvement in inhibitory performance during childhood and adolescence, but this transition is likely aided by child-unique mechanisms.

This study cannot give a causal explanation for the neural and behavioral changes during development. However, this is a comprehensive study that contrasted source models of adult and child auditory processing and investigated the relevance for response inhibition. The experiment was carefully designed to limit external factors (e.g., differences in motor and/or stimuli between conditions). Thus, this is the most direct comparison possible between active and passive tasks. In combination with our methods (combined M/EEG and individual MRIs) it provides greater confidence in our conclusion that associations between auditory activation and inhibition task performance differ in adults and children.

Future studies should look at individual changes over time (i.e., longitudinal) to investigate whether the developmental changes relate to the improved performance in response inhibition. Ideally, this should also include magnetic resonance spectroscopy (MRS) to measure the biochemical changes in the brain, as maturation of GABAergic versus glutamatergic circuits likely play a crucial role in both auditory and response inhibition development (Du et al., 2018; Le Magueresse & Monyer, 2013; Sanes & Kotak, 2011; Silveri et al., 2013). Moreover, an important remaining question is whether the effects are limited to the left hemisphere or depend on handedness.

The clinical importance of achieving competent response inhibition highlights the value of a complete understanding of the typical development of this process. This study shows remarkable differences between adults and children in (auditory) processing during response inhibition. It emphasizes the cruciality of sensory processing during (critical) periods of development until adult-like response inhibition networks have matured sufficiently. Future studies that examine the source models longitudinally, and account for biochemical changes, would be welcomed.

## ACKNOWLEDGMENTS

We are grateful to Hanna-Majja Lapinkero, Suvi Karjalainen, Maria Vesterinen & Janne Rajaniemi for help with data collection and to Amit Jaiswal, Erka Heinilä and Jukka Nenonen for their help with preprocessing and scripting. This work was supported by EU project ChildBrain (Horizon2020 Marie Skłodowska-Curie Action (MSCA) Innovative Training Network (ITN)—European Training Network (ETN), grant agreement no. 641652) and the Academy of Finland grant number 311877.

## CONFLICT OF INTEREST STATEMENT

None.

## DATA AVAILABILITY STATEMENT

Raw data are not publicly available due to legal restrictions. Derived data supporting the findings of this study are available here: <https://osf.io/rhb5z/>.

## ORCID

Sam van Bijnen  <https://orcid.org/0000-0003-0023-6032>

Tiina Parviainen  <https://orcid.org/0000-0001-6992-5157>

## REFERENCES

- Albrecht, R., Suchodoletz, W. V., & Uwer, R. (2000). The development of auditory evoked dipole source activity from childhood to adulthood. *Clinical Neurophysiology*, 111(12), 2268–2276.
- Baillet, S. (2017). Magnetoencephalography for brain electrophysiology and imaging. *Nature Neuroscience*, 20(3), 327–339.
- Benner, J., Wengenroth, M., Reinhardt, J., Stippich, C., Schneider, P., & Blatow, M. (2017). Prevalence and function of Heschl's gyrus morphotypes in musicians. *Brain Structure and Function*, 222(8), 3587–3603.
- Botvinick, M. M., Cohen, J. D., & Carter, C. S. (2004). Conflict monitoring and anterior cingulate cortex: An update. *Trends in Cognitive Sciences*, 8(12), 539–546.
- Čeponienė, R., Rinne, T., & Näätänen, R. (2002). Maturation of cortical sound processing as indexed by event-related potentials. *Clinical Neurophysiology*, 113(6), 870–882.

- Chambers, C. D., Garavan, H., & Bellgrove, M. A. (2009). Insights into the neural basis of response inhibition from cognitive and clinical neuroscience. *Neuroscience & Biobehavioral Reviews*, 33(5), 631–646.
- Dale, A. M., Fischl, B., & Sereno, M. I. (1999). Cortical surface-based analysis: I. Segmentation and surface reconstruction. *Neuroimage*, 9(2), 179–194.
- Desikan, R. S., Ségonne, F., Fischl, B., Quinn, B. T., Dickerson, B. C., Blacker, D., Buckner, R. L., Dale, A. M., Maguire, R. P., Hyman, B. T., Albert, M. S., & Killiany, R. J. (2006). An automated labeling system for subdividing the human cerebral cortex on MRI scans into gyral based regions of interest. *NeuroImage*, 31(3), 968–980.
- Du, X., Rowland, L. M., Summerfelt, A., Wijtenburg, A., Chiappelli, J., Wisner, K., Kochunov, P., Choa, F.-S., & Hong, L. E. (2018). TMS evoked N100 reflects local GABA and glutamate balance. *Brain Stimulation*, 11(5), 1071–1079.
- Eggermont, J. J., & Ponton, C. W. (2003). Auditory-evoked potential studies of cortical maturation in normal hearing and implanted children: Correlations with changes in structure and speech perception. *Acta Oto-Laryngologica*, 123(2), 249–252.
- Engemann, D. A., & Gramfort, A. (2015). Automated model selection in covariance estimation and spatial whitening of MEG and EEG signals. *NeuroImage*, 108, 328–342.
- Falkenstein, M., Hoormann, J., & Hohnsbein, J. (1999). ERP components in Go/Nogo tasks and their relation to inhibition. *Acta Psychologica*, 101(2–3), 267–291.
- Fischl, B., Sereno, M. I., & Dale, A. M. (1999). Cortical surface-based analysis II: Inflation, flattening, and a surface-based coordinate system. *NeuroImage*, 9, 195–207.
- Fischl, B., Sereno, M. I., Tootell, R. B., & Dale, A. M. (1999). High-resolution intersubject averaging and a coordinate system for the cortical surface. *Human Brain Mapping*, 8, 272–284.
- Gramfort, A., Luessi, M., Larson, E., Engemann, D. A., Strohmeier, D., Brodbeck, C., Goj, R., Jas, M., Brooks, T., & Hämäläinen, M. (2013). MEG and EEG data analysis with MNE-Python. *Frontiers in Neuroscience*, 267, 1–13.
- Gramfort, A., Luessi, M., Larson, E., Engemann, D. A., Strohmeier, D., Brodbeck, C., Parkkonen, L., & Hämäläinen, M. S. (2014). MNE software for processing MEG and EEG data. *NeuroImage*, 86, 446–460.
- Hämäläinen, M. S., & Ilmoniemi, R. J. (1994). Interpreting magnetic fields of the brain: Minimum norm estimates. *Medical & Biological Engineering & Computing*, 32(1), 35–42.
- Hämäläinen, J. A., Salminen, H. K., & Leppänen, P. H. (2013). Basic auditory processing deficits in dyslexia: Systematic review of the behavioral and event-related potential/field evidence. *Journal of Learning Disabilities*, 46(5), 413–427.
- Huster, R. J., Westerhausen, R., Pantev, C., & Konrad, C. (2010). The role of the cingulate cortex as neural generator of the N200 and P300 in a tactile response inhibition task. *Human Brain Mapping*, 31(8), 1260–1271.
- Hyvärinen, A., & Oja, E. (2000). Independent component analysis: Algorithms and applications. *Neural Networks*, 13(4–5), 411–430.
- Jas, M., Engemann, D. A., Bekhti, Y., Raimondo, F., & Gramfort, A. (2017). Autoreject: Automated artifact rejection for MEG and EEG data. *NeuroImage*, 159, 417–429.
- Johnstone, S. J., Barry, R. J., Anderson, J. W., & Coyle, S. F. (1996). Age-related changes in child and adolescent event-related potential component morphology, amplitude and latency to standard and target stimuli in an auditory oddball task. *International Journal of Psychophysiology*, 24(3), 223–238.
- Kohl, C., Parviainen, T., & Jones, S. R. (2022). Neural mechanisms underlying human auditory evoked responses revealed by human neocortical neurosolver. *Brain Topography*, 35(1), 19–35.
- Le Magueresse, C., & Monyer, H. (2013). GABAergic interneurons shape the functional maturation of the cortex. *Neuron*, 77(3), 388–405.
- Muthén, L. K., & Muthén, B. O. (2012). *Mplus user's guide* (7th ed.). Muthén & Muthén.
- Moore, J. K., & Guan, Y. L. (2001). Cytoarchitectural and axonal maturation in human auditory cortex. *Journal of the Association for Research in Otolaryngology*, 2(4), 297–311.
- Moore, J. K., & Linthicum, F. H., Jr. (2007). The human auditory system: A timeline of development. *International Journal of Audiology*, 46(9), 460–478.
- Näätänen, R. (1990). The role of attention in auditory information processing as revealed by event-related potentials and other brain measures of cognitive function. *Behavioral and Brain Sciences*, 13(2), 201–233.
- Nieuwenhuis, S., Yeung, N., Van Den Wildenberg, W., & Ridderinkhof, K. R. (2003). Electrophysiological correlates of anterior cingulate function in a go/no-go task: Effects of response conflict and trial type frequency. *Cognitive, Affective, & Behavioral Neuroscience*, 3(1), 17–26.
- Orehkova, E. V., Butorina, A. V., Tsetlin, M. M., Novikova, S. I., Sokolov, P. A., Elam, M., & Stroganova, T. A. (2013). Auditory magnetic response to clicks in children and adults: Its components, hemispheric lateralization and repetition suppression effect. *Brain Topography*, 26(3), 410–427.
- Orehkova, E. V., Tsetlin, M. M., Butorina, A. V., Novikova, S. I., Gratchev, V. V., Sokolov, P. A., Elam, M., & Stroganova, T. A. (2012). Auditory cortex responses to clicks and sensory modulation difficulties in children with autism spectrum disorders (ASD). *PLoS One*, 7(6), e39906.
- Paetau, R., Ahonen, A., Salonen, O., & Sams, M. (1995). Auditory evoked magnetic fields to tones and pseudowords in healthy children and adults. *Journal of Clinical Neurophysiology*, 12(2), 177–185.
- Parviainen, T., Helenius, P., Poskiparta, E., Niemi, P., & Salmelin, R. (2011). Speech perception in the child brain: Cortical timing and its relevance to literacy acquisition. *Human Brain Mapping*, 32(12), 2193–2206.
- Parviainen, T., Helenius, P., & Salmelin, R. (2019). Children show hemispheric differences in the basic auditory response properties. *Human Brain Mapping*, 40(9), 2699–2710.
- Parviainen, T., Helenius, P., & Salmelin, R. (2005). Cortical differentiation of speech and nonspeech sounds at 100 ms: Implications for dyslexia. *Cerebral Cortex*, 15(7), 1054–1063.
- Peirce, J., Gray, J. R., Simpson, S., MacAskill, M., Höchenberger, R., Sogo, H., Kastman, E., & Lindeløv, J. K. (2019). PsychoPy2: Experiments in behavior made easy. *Behavior Research Methods*, 51(1), 195–203.
- Piastra, M. C., Nüßing, A., Vorwerk, J., Clerc, M., Engwer, C., & Wolters, C. H. (2021). A comprehensive study on electroencephalography and magnetoencephalography sensitivity to cortical and subcortical sources. *Human Brain Mapping*, 42(4), 978–992.
- Picton, T. W., Hillyard, S. A., Krausz, H. I., & Galambos, R. (1974). Human auditory evoked potentials. I: Evaluation of components. *Electroencephalography and Clinical Neurophysiology*, 36, 179–190.
- Ponton, C., Eggermont, J. J., Khosla, D., Kwong, B., & Don, M. (2002). Maturation of human central auditory system activity: Separating auditory evoked potentials by dipole source modeling. *Clinical Neurophysiology*, 113(3), 407–420.
- Ponton, C. W., Eggermont, J. J., Kwong, B., & Don, M. (2000). Maturation of human central auditory system activity: Evidence from multi-channel evoked potentials. *Clinical Neurophysiology*, 111(2), 220–236.
- Puii, A. A., Wudarczyk, O., Kohls, G., Bzdok, D., Herpertz-Dahlmann, B., & Konrad, K. (2020). Meta-analytic evidence for a joint neural mechanism underlying response inhibition and state anger. *Human Brain Mapping*, 41(11), 3147–3160.
- Ruhnau, P., Herrmann, B., Maess, B., & Schröger, E. (2011). Maturation of obligatory auditory responses and their neural sources: Evidence from EEG and MEG. *NeuroImage*, 58(2), 630–639.
- Sanes, D. H., & Kotak, V. C. (2011). Developmental plasticity of auditory cortical inhibitory synapses. *Hearing Research*, 279(1–2), 140–148.

- Shvarts, V., & Mäkelä, J. P. (2020). Auditory mapping with MEG: An update on the current state of clinical research and practice with considerations for clinical practice guidelines. *Journal of Clinical Neurophysiology*, 37(6), 574–584.
- Shahin, A. J., Roberts, L. E., Miller, L. M., McDonald, K. L., & Alain, C. (2007). Sensitivity of EEG and MEG to the N1 and P2 auditory evoked responses modulated by spectral complexity of sounds. *Brain Topography*, 20(2), 55–61.
- Shaw, M. E., Hämäläinen, M. S., & Gutschalk, A. (2013). How anatomical asymmetry of human auditory cortex can lead to a rightward bias in auditory evoked fields. *NeuroImage*, 74, 22–29.
- Silveri, M. M., Sneider, J. T., Crowley, D. J., Covell, M. J., Acharya, D., Rosso, I. M., & Jensen, J. E. (2013). Frontal lobe  $\gamma$ -aminobutyric acid levels during adolescence: Associations with impulsivity and response inhibition. *Biological Psychiatry*, 74(4), 296–304.
- Smith, J. L., Johnstone, S. J., & Barry, R. J. (2007). Response priming in the Go/NoGo task: The N2 reflects neither inhibition nor conflict. *Clinical Neurophysiology*, 118(2), 343–355.
- Stuss, D. T., Murphy, K. J., Binns, M. A., & Alexander, M. P. (2003). Staying on the job: The frontal lobes control individual performance variability. *Brain*, 126(11), 2363–2380.
- Sussman, E., Steinschneider, M., Gumenyuk, V., Grushko, J., & Lawson, K. (2008). The maturation of human evoked brain potentials to sounds presented at different stimulus rates. *Hearing Research*, 236(1–2), 61–79.
- Takeshita, K., Nagamine, T., Thuy, D. H. D., Satow, T., Matsuhashi, M., Yamamoto, J., Takayama, M., Fujiwara, N., & Shibasaki, H. (2002). Maturation change of parallel auditory processing in school-aged children revealed by simultaneous recording of magnetic and electric cortical responses. *Clinical Neurophysiology*, 113(9), 1470–1484.
- van Bijnen, S., Kärkkäinen, S., Helenius, P., & Parviainen, T. (2019). Left hemisphere enhancement of auditory activation in language impaired children. *Scientific Reports*, 9(1), 1–11.
- van Bijnen, S., Parkkonen, L., & Parviainen, T. (2022). Activity level in left auditory cortex predicts behavioral performance in inhibition tasks in children. *NeuroImage*, 258, 119371.
- Wechsler, D. (1999). *WISC-III Käsikirja [manual for the Wechsler intelligence scale for children—Third edition. Manual for the Finnish adaptation]*. Psykologien kustannus Oy.
- Weiss, H., & Luciana, M. (2022). Neurobehavioral maturation of motor response inhibition in adolescence—A narrative review. *Neuroscience & Biobehavioral Reviews*, 137, 104646.
- Wunderlich, J. L., Cone-Wesson, B. K., & Shepherd, R. (2006). Maturation of the cortical auditory evoked potential in infants and young children. *Hearing Research*, 212(1–2), 185–202.
- Yoshimura, Y., Kikuchi, M., Ueno, S., Shitamichi, K., Remijn, G. B., Hiraishi, H., Hasegawa, C., Furutani, N., Oi, M., Munesue, T., Tsubokawa, T., Higashida, H., & Minabe, Y. (2014). A longitudinal study of auditory evoked field and language development in young children. *NeuroImage*, 101, 440–447.

### SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article.

**How to cite this article:** van Bijnen, S., Muotka, J., & Parviainen, T. (2023). Divergent auditory activation in relation to inhibition task performance in children and adults. *Human Brain Mapping*, 1–14. <https://doi.org/10.1002/hbm.26418>